## Patent Claims

- Stable formulation of a solution consisting of xylometazoline hydrochloride and oxymetazoline hydrochloride as active substance, a solvent which is pharmacologically acceptable for nasal administration, an adjuvant selected from among sorbitol and/or glycerol and an inorganic pH buffer.
- 10 2. Formulation according to claim 1, characterised in that the active substance is present in a concentration of between 0.01 and 1.0% by weight, preferably between 0.01 and 0.5% by weight and most preferably between 0.05 and 0.1% by weight.
  - Formulation according to one of claims 1 and 2, characterised in that the solvent is water.
- 4. Formulation according to one of claims 1 and 2, characterised in that the solvent is a mixture of ethanol and water.
- 5. Formulation according to one of claims 1 to 4, characterised in that the proportion of adjuvant in the solution is 1 to 10% by weight, preferably 2 to 6% by weight.
  - 6. Formulation according to claim 5, characterised in that the adjuvant is 3.5 to 4.5% by weight, preferably 4.0% by weight, of sorbitol.
    - 7. Formulation according to claim 6, characterised in that the adjuvant is 2.0 to 2.8% by weight, preferably 2.4% by weight, of glycerol.

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8. Formulation according to one of claims 1 to 7, characterised in that the solution contains a sodium and/or potassium phosphate buffer or a sodium and/or potassium borate buffer.

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9. Formulation according to one of claims 1 to 8, characterised in that the solution contains a monosodium dihydrogen-disodium monohydrogen phosphate buffer and/or monopotassium dihydrogen-dipotassium monohydrogen phosphate buffer.

10. Formulation according to one of claims 1 to 9, characterised in that the solution is adjusted to a pH of 5.0 to 6.8, preferably 5.5 to 6.8, most preferably 5.8 to 6.0.

11. Formulation according to one of claims 1 to 10, characterised in that the formulation contains an oligodynamically active substance.

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- 12. Formulation according to claim 11, characterised in that the oligodynamic substance is silver or silver ions.
- 25 13. Formulation according to one of claims 1 to 12, characterised in that the formulation contains only xylometazoline hydrochloride as active substance.
- 14. Formulation according to one of claims 1 to 12,
  30 characterised in that the formulation contains only oxymetazoline hydrochloride as active substance.
- 15. Use of a formulation according to claims 1 to 14 together with an inhaler having silver-containing elements in the region between the active substance reservoir and the sprayhead.

16. Use of the formulation according to one of claims 1 to 13 as a rhinological agent.